

## REVIEW ARTICLE

# Total Ankle Replacement

Indications, Implant Designs, and Results

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## SUMMARY

**Background:** About 1% of adults suffer from painful osteoarthritis of the ankle. The current literature contains no information on the percentage of such patients who derive long-term relief of symptoms from conservative treatment. Advanced ankle osteoarthritis can be treated with non-joint-preserving measures, such as total ankle replacement and ankle fusion.

**Methods:** This review is based on selected relevant publications, guidelines from Germany and abroad, and the authors' personal experience.

**Results:** Before surgery is considered, conservative measures such as physiotherapy and orthopedic aids should be used to the fullest possible extent. No randomized trials have yet been published comparing total ankle replacement with ankle fusion. Total ankle replacement with newer types of prosthesis yields good to very good intermediate-term and long-term results, with mean success rates of up to 90% at 10 years (range, 68–100%). Independent risk factors for the failure of ankle replacement are age over 70 years (odds ratio [OR] 3.84), primary osteoarthritis (OR 7.19), post-traumatic osteoarthritis (OR 6.2), and type of prosthesis (e.g., single hydroxyapatite coating: OR 15.04). The average range of motion of the replaced ankle joint is 25° to 30°, with values as high as 60°.

**Conclusion:** Total ankle replacement is a good treatment option for complete, end-stage ankle arthritis. It can restore joint function and make the patient mobile with little or no pain. There are, however, many contraindications to be taken into account. There is a need for further studies of the biomechanics of arthritic and replaced ankle joints and for long-term follow-up studies of total ankle replacement.

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Osteoarthritis of the ankle is an increasing issue in the healthcare sector (1, e1, e2). Approximately 1% of the adult population suffers from painful ankle osteoarthritis (2). The psychological and physical limitations associated with advanced ankle osteoarthritis are at least as marked as those of patients with osteoarthritis of the knee or hip (2). Degenerative changes of the ankle, in contrast to osteoarthritis of the knee or hip, are usually posttraumatic (*Table 1, eTable 1*) (3, e3, e4). Both poorly healed fractures to the lower extremity (4, e5) and repetitive ligament injuries (5) can play a major role. The main causes of secondary osteoarthritis of the ankle include rheumatic diseases, hemophilia, hemochromatosis, gout, avascular necrosis, and postinfectious states (1, e1).

This review article uses the current literature to explain the indications and the absolute and relative contraindications for total ankle replacement. It also presents the results of current clinical studies on post-operative functional outcomes and the probability of success of ankle replacement surgery.

## Selective literature search

This review article is based on a selective literature search in established databases. The following medical databases were searched, with no date restriction: Medline, Cochrane, EmbaseTM, Cinahl, Google Scholar, ScienceDirect, and SpringerLink. The search terms used were the following: “total ankle replacement,” “total ankle arthroplasty,” “ankle replacement,” “ankle arthroplasty,” and “ankle prosthesis.” All articles written in languages spoken by the authors (German, English, and French) were included.

The digital indices of the following orthopedic journals were also searched for the above-mentioned search terms: *Foot and Ankle International*; *Journal of Bone and Joint Surgery, American Volume*; *Bone & Joint Journal* (formerly known as the *Journal of Bone and Joint Surgery, British Volume*); *Clinical Orthopaedics and Related Research*; *Foot and Ankle Clinics of North America*; *Journal of Foot and Ankle Surgery*; and *Der Orthopäde*. In addition, the bibliographies of the identified original and review articles were searched for further studies.

The literature search was performed by two of the authors (AB and MDW), independently of each other.

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**TABLE 1**

**Etiology of advanced ankle osteoarthritis, based on a selection of clinical and epidemiological studies\***

Study	Study type	Patients (ankles)	Etiology of ankle osteoarthritis, % (absolute values)		
			Primary	Secondary	Posttraumatic
(e6)	RS, SC, clinical	45 (51)	25.5% (13)	54.9% (28)	19.6% (10)
(6)	PS, SC, clinical	684 (722)	9.5% (69)	11.4% (82)	79.1% (571)
(e7)	PS, SC, clinical	47 (50)	6.0% (3)	8.0% (4)	86.0% (43)
(e8)	PS, SC, clinical	49 (50)	16.0% (8)	18.0% (9)	66.0% (33)
(e9)	PS, MC register, clinical	245 (257)	20.6% (53)	55.3% (142)	24.1% (62)
(e10)	PS, SC, clinical	80 (83)	33.7% (28)	25.3% (21)	41.0% (34)
(e11)	RS, SC, clinical	111 (123)	52.8% (65)	18.7% (23)	28.5% (35)
(e12)	RS, SC, clinical	61 (62)	19.4% (12)	4.8% (3)	75.8% (47)
(e13)	RS, SC, clinical	45 (52)	50.0% (26)	26.9% (14)	23.1% (12)
(e14)	RS, SC, clinical	126 (132)	46.2% (61)	25.0 (33)	28.8% (38)
(e15)	PS, SC, clinical	43 (50)	54.0% (27)	32.0% (16)	14.0% (7)
(e16)	PS, SC, clinical	396 (404)	16.6% (67)	13.6% (55)	69.8% (282)
(e17)	PS, SC, clinical	80 (84)	25.0% (21)	19.0% (16)	56.0% (47)
(e18)	PS, SC, clinical	82 (82)	34.2% (28)	13.4% (11)	52.4% (43)
(e19)	RS, SC, clinical	95 (100)	26.0% (26)	29.0% (29)	45.0% (45)
(e20)	PS, SC, clinical	229 (229)	13.8% (32)	4.0% (9)	82.2% (188)
(e21)	PS, SC, clinical	106 (106)	52.8% (56)	20.8% (22)	26.4% (28)
(e22)	PS, SC, clinical	233 (240)	30.8% (74)	17.9% (43)	51.3% (123)
(3)	PS, SC, epidemiological	639 (639)	7.2% (46)	23.2% (148)	69.6% (445)
(7)	PS, MC, clinical	593 (593)	26.5% (157)	15.3% (91)	58.2% (345)
(e23)	RS, SC, clinical	100 (100)	30.0% (30)	44.0% (44)	26.0% (26)
(e24)	RS, MC, clinical	501 (517)	13.9% (72)	25.9% (134)	60.2% (311)
(e25)	PS, MC register, clinical	515 (515)	19.2% (99)	59.2% (305)	21.6% (111)
(e26)	RS, SC, clinical	303 (306)	25.2% (77)	10.1% (31)	64.7% (198)
(e27)	RS, SC, clinical	103 (103)	71.8% (74)	18.5% (19)	9.7% (10)
(e28)	PS, SC, clinical	65 (68)	13.2% (9)	16.2% (11)	70.6% (48)
(e3)	RS, SC, epidemiological	390 (406)	8.9% (36)	12.8% (52)	78.3% (318)
(e29)	PS, SC, clinical	66 (66)	0.0% (0)	10.6% (7)	89.4% (59)
(e4)	RS, SC, clinical	226 (233)	5.6% (13)	23.2% (54)	71.2% (166)
(e30)	PS, SC, clinical	96 (100)	64.0% (64)	27.0% (27)	9.0% (9)
(e31)	RS, SC, clinical	90 (99)	40.4% (40)	12.1% (12)	47.5% (47)
Total/mean		6218 (6389)	21.5% (1373)	22.6% (1373)	56.0% (3575)

\*Clinical (total ankle replacement) and epidemiological (etiology of ankle osteoarthritis) studies with at least 50 patients were included. MC, multicenter; PS, prospective; RS, retrospective; SC, single-center

**History and implant designs**

Most first-generation ankle replacements performed in the 1970s and early 1980s were two-component cemented implants. The rate of aseptic loosening for all first-generation implant types was extremely high, occurring in almost 90% of implants (8).

Second-generation ankle implants (from the mid-1980s onwards) show improved implant shapes and better surgical technique: bone-conserving

surgical approach and no cementation. Today there are several commercially available ankle implant types (*Figure 1*). All implant designs can be classified by surgical technique and implant properties (*eTable 2*) (8).

**Diagnosis and preoperative planning**

A clinical and radiological diagnosis of osteoarthritis of the ankle can be made by the patient's treating

physician on the basis of clinical and radiological examination, as described.

The first step in preoperative diagnosis is to take a clinical history. All available documents should be evaluated: it is important to note which, if any, treatment options have already been administered. Further information such as BMI (body mass index), physical activity levels, previous and/or current treatment, severity of pain, limitations in everyday private and/or occupational activities, intake of analgesics, and concomitant diagnoses (diabetes mellitus, osteoporosis, polyneuropathy, etc.) should be recorded.

Clinical examination begins with examination of the foot/hindfoot on standing, sitting, and walking. Hind-foot alignment (valgus, varus, or neutral) is assessed from behind, with the patient standing. Stability is determined with the patient seated, using the talar tilt test (examination of medial and lateral ankle inversion) and the anterior drawer test (which tests for increased anterior translation of the talus) (9). Mobility of the subtalar joint is measured using a goniometer under load (10). Mobility of the ankle is measured manually, with the ankle fixed and free (e32).

Radiological examination includes conventional weight bearing radiographs: dorsoplantar and lateral views of the foot, anteroposterior (mortise) and lateral views of the ankle, and Saltzman view (hindfoot alignment view to assess inframalleolar alignment [11]) (Figure 2). Supramalleolar alignment is determined using the medial distal tibial angle (12, e33). In patients with knee deformities, a whole leg radiograph (orthoradiogram) is also taken. Optionally, computed tomography or magnetic resonance imaging may be performed; these can provide important additional information.

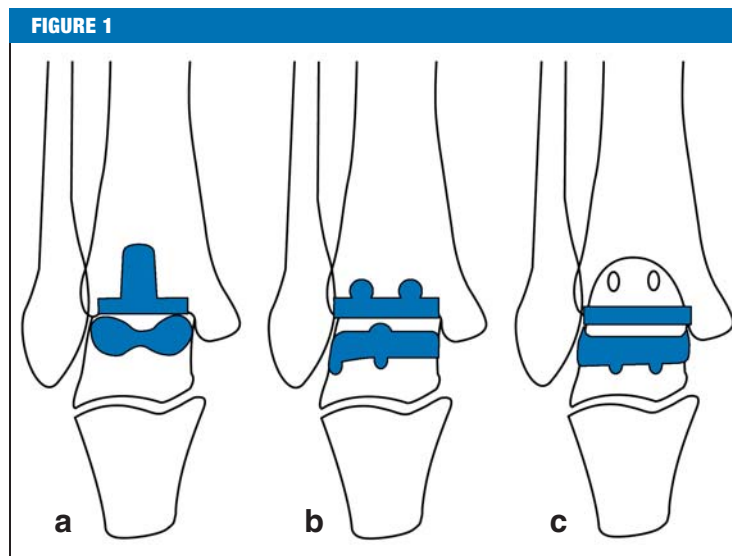
### Indication for surgery

Conservative therapy should be administered before surgery is indicated. This includes intensive physiotherapy (local antiphlogistic therapy, muscle and movement exercises to prevent stiffness of the joint, muscle strength development, gait training) and possibly intra-articular hyaluronic acid viscosupplementation and orthopedic adaptation of footwear (13, 14).

The ideal indication for total ankle replacement is advanced, complete osteoarthritis of the ankle (primary, secondary, or posttraumatic) with good bone quality, neutral alignment, good stability, and preserved mobility of the ankle. Further special indications include patients with bilateral osteoarthritis of the ankle (15, e34).

Total joint replacement can also be performed as revision arthroplasty in patients with failed ankle prosthesis (16, 17, e35). However, revision ankle arthroplasty, like revision joint replacement in general, is a technically demanding surgical procedure. Patients with painful non-union or malunion of previous ankle arthrodesis are another specific indication for total ankle replacement (18, e36, e37).

Absolute contraindications include acute or chronic infections, with or without osteomyelitis or osteitis;



**Modern ankle implant types**

a) Components with tibial stem b) Components with bars c) Flat components

severe osteonecrosis of the talus (more than one third of the talus); neuromuscular diseases; neuroarthropathies (e.g. patients with Charcot foot); and patients with severe circulatory disorders (19). In patients with concomitant significant ligament instabilities and/or deformities that cannot be corrected intraoperatively, arthrodesis of the ankle should be performed instead of joint replacement. Metal allergies are also a contraindication (20, 21).

Relative contraindications include severe osteoporosis, poor bone quality (e.g. due to steroid treatment), diabetes mellitus, smoking, and excess weight, although the literature shows that good outcomes can be achieved in some of these cases [22]). There may be an increased rate of aseptic loosening of implant components in patients who engage in high levels of sporting activity (23, 24). Low-impact exercise (walking, swimming, cycling, golf), however, is recommended postoperatively (19, 24).

### Surgical technique

An anterior approach is usually used for ankle replacement surgery (eFigure 1). In patients with a history of previous ankle surgery, the surgical approach can be modified in order not to compromise postoperative wound healing (e38, e39). Depth preparation is performed beneath the tendon of the tibialis anterior muscle in order to preserve the anterior neurovascular bundle, which in most cases lies behind the tendon of the extensor hallucis longus muscle or between the tendons of the extensor hallucis longus and extensor digitorum longus muscles (e40). Bone resection is performed using an oscillating saw. Additional procedures for patients with concomitant deformities and/or instabilities should be performed after insertion of the implant components (eTable 3) (25, 26, e41, e42).



**Figure 2:** Preoperative conventional X-ray in standing position of 67-year-old female patient with posttraumatic ankle osteoarthritis following open reduction and internation fixation for trimalleolar luxation fracture 4 years earlier: a) mortise view of ankle; b) lateral view of foot and ankle; c) dorsoplantar view of foot; d) Saltzman view of hindfoot

**Aftercare**

We recommend immobilization using plaster cast of the lower leg or a stabilizing boot for six weeks after surgery. During this period full weight may be borne with the aid of two elbow crutches, depending on the severity of the patient’s complaints. In patients with reduced bone quality and/or who have undergone additional procedures such as corrective osteotomy, we recommend 15 kg partial weightbearing for six weeks after surgery. Thromboprophylaxis is administered during immobilization (27). Clinical and radiological follow-up examination is performed after six weeks (Figure 3). After this, intensive outpatient physiotherapy begins: gait training, proprioception exercises, gradual increase to full weightbearing, local antiphlogistic therapy including lymph drainage, active and passive ankle mobility therapy, extension exercises, and therapy to strengthen the triceps surae muscle.

Compression stockings are used for patients with persistent edema or soft-tissue swelling. The following sports can be recommended after full mobilization and full weightbearing ability have been attained: low-impact (e.g. walking, swimming, cycling, golf) or medium-impact (e.g. jogging, tennis, skiing) (24). Contact sports and sports that involve jumping should be avoided (24).

Clinical and radiological follow-up examinations are performed six weeks, three months, six months, and one year after surgery and then annually. The most important tools/questionnaires (28) that can be used to record functional postoperative outcomes following total ankle replacement are the American Orthopaedic Foot and Ankle Society (AOFAS) Ankle–Hindfoot Score (29) (score composed of pain, function, and alignment; minimum score, 0 points; maximum score, 100 points); and the Kofoed Ankle Score (e58) (score

composed of pain, function, and range of motion; minimum score, 0 points; maximum score, 100 points). Pain level is determined using the visual analogue scale (VAS) from 0 (no pain) to 10 (worst possible pain) (e59). Quality of life can be analyzed using the SF-36 questionnaire (36-Item Short Form Health Survey) (e60).

**Results/literature review**

For a long time arthrodesis of the ankle was the first-line treatment for patients with advanced osteoarthritis of the ankle, which is not surprising given how uniformly disappointing the results of first-generation total ankle replacement were. Precise analysis of failures led to the development of new implant designs, acceptance of which is steadily increasing among orthopedic surgeons.

However, it is difficult to find well conducted, controlled, prospective studies in the literature, and in particular there are no comparisons of two-component and three-component implant types (30). Saltzman et al. (7) published the first results of a prospective study comparing ankle arthrodesis (66 cases) and total ankle replacement (593 patients) and demonstrated that patients with total ankle replacement had less pain and better functional outcomes postoperatively, with comparable postoperative complication rates. Although postoperative complications (poor wound healing, infections) were observed more frequently in patients undergoing ankle replacement than in those undergoing arthrodesis of the ankle—6.2% versus 1.5%—the difference was not statistically significant ( $p = 0.087$ ). The Buechel–Pappas score (score composed of pain, function, deformity, and mobility; minimum score, 0 points; maximum score, 100 points) (e61) was used to assess functional outcome. Patients with total ankle

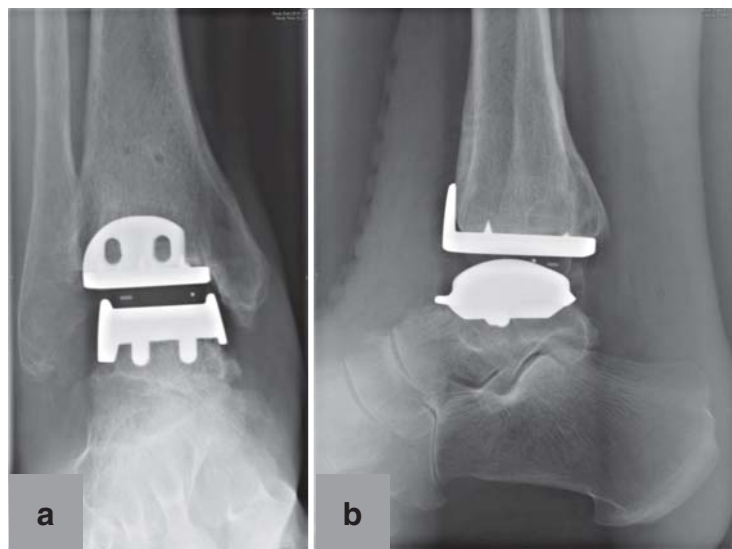


replacement had significantly better functional outcomes: Buechel–Pappas score  $46.7 \pm 13.0$  versus  $26.3 \pm 17$  ( $p < 0.001$ ). The two groups had comparable postoperative pain levels:  $1.6 \pm 1.8$  versus  $1.8 \pm 2.0$  ( $p = 0.607$ ). Further studies are planned by the authors but have not yet been published (7).

Despite increasing acceptance, total ankle replacement remains a technically demanding procedure with a flat learning curve. Intraoperative complications are not uncommon; they include fractures of the medial and/or lateral malleolus in 0 to 23% of cases and tendon injuries (posterior tibial tendon, flexor hallucis longus, flexor digitorum longus) and nerve injuries (superficial/deep peroneal nerve) in 0 to 10% of cases (31, e62–e66). Difficult steps during surgery include correct component positioning, particularly of talar components (e62, e66). Incorrect tibial component positioning can be found in 0 to 16% of all cases, and incorrect talar component positioning in 0 to 36% of all cases (e62). Numerous *in vitro* biomechanical studies have shown that incorrect positioning of implant components has adverse biomechanical consequences such as reduced ankle mobility, pathological tension of the periarticular ligaments, and unfavorable intra-articular pressure distribution (e67–e70). We have shown in a clinical study that patients with suboptimal positioning of talar components have a higher rate of persisting pain and worse ankle mobility (32).

Postoperative outcomes following total ankle replacement are steadily improving (Table 2; eTable 4) but lag behind those of total knee and hip replacements (Table 3). Labek et al. (33) investigated cumulative outcomes on the basis of worldwide joint replacement registers. Outcomes following total hip and knee replacements were comparable, with 1.29 and 1.26 revisions per 100 component years. This means that after 10 years 13 out of every 100 patients need to undergo revision surgery. The outcomes following medial partial replacement were somewhat worse, with 1.53 revisions per 100 component years. Total ankle replacement was associated with the worst outcomes, however, with 3.29 revisions per 100 component years, resulting in revision surgery for 33 out of every 100 patients within 10 years (33). The causes and frequency of failure of total ankle replacement are different from those of hip and knee replacements: the main causes of failure are aseptic loosening of tibial and/or talar components, persisting pain, and septic loosening (Table 3) (34).

In 2010, Gougoulias et al. (35) performed a systematic review of the literature including 13 level IV studies with a total of 1105 ankle replacements. Seven different implant types were used. The mean failure rate (defined as replacement of one or both implant components or implant removal and conversion to arthrodesis of the ankle) five years after implantation was 10%, but there was great variation in failure rates between different centers, ranging from 0% to 32%. The percentage of patients in the included studies with persisting complaints was between 27 and 60%.



**Figure 3:** Postoperative X-ray of 67-year-old female patient 6 weeks after total ankle replacement: a) mortise view of ankle; b) lateral view of ankle

Postoperative improvement in ankle mobility was relatively low, with values between  $0^\circ$  and  $14^\circ$  (35). Zaidi et al. (36) published a systematic review of the literature and meta-analysis of 58 publications with a total of 7942 ankle replacements. The success rate after 10 years was 89%, with an annual failure rate of 1.2% (95% confidence interval [CI]: 0.7 to 1.6). The mean AOFAS Ankle–Hindfoot Score rose from 40 (95% CI: 36 to 43) preoperatively to 80 (95% CI: 76 to 84) postoperatively. The range of motion of the ankle on which surgery was performed improved from a mean of  $23^\circ$  (95% CI:  $19$  to  $26^\circ$ ) preoperatively to  $34^\circ$  (95% CI:  $26$  to  $41^\circ$ ) postoperatively (36).

We performed a survivorship analysis of implant components in 684 patients who received a total of 722 ankle replacements (6). The mean follow-up time in this prospective study was  $6.3 \pm 2.9$  years. The probability of success of the implant components was 94% after five years and 84% after 10 years. These results are comparable with those of current clinical studies (Table 2, eTable 4). The following factors were identified as independent risk factors for ankle replacement failure:

- Age under 70 years (odds ratio [OR]: 3.84)
- Etiology of ankle osteoarthritis (OR for primary osteoarthritis: 7.19; OR for posttraumatic osteoarthritis: 6.20)
- Implant generation (OR for single hydroxyapatite coating: OR: 15.04) (6).

For a long time a change of approach—removal of the implant components followed by arthrodesis—was the standard procedure in cases of ankle replacement failure. The current literature describes various surgical techniques and fixation methods for such arthrodesis after prosthesis removal: bone allografts, autografts, or replacement materials (e.g. porous metals such as

**TABLE 2**

**Clinical outcomes following total ankle replacement: probability of survival of implant components\***

Study	Study type	Implant type	No. of implants	Probability of survival of implant components
(e72)	RS, SC	Agility	207	76% after 9 years
(22)	RS, SC	HINTEGRA	123	93% after 6 years
(6)	PS, SC	HINTEGRA	722	94% after 5 years, 84% after 10 years
(e74)	RS, MC	Salto (388), AES (173), HINTEGRA (22), STAR (9)	592	88% after 71 months
(e79)	PS, SC	Buechel-Pappas (normal sulcus 40; deep sulcus 75)	115	74.2% (normal sulcus) after 20 years, 92% (deep sulcus) after 12 years
(e9)	PS, MC	STAR (216), TPR (32), HINTEGRA (6), AES (3)	257	89% after 5 years, 76% after 10 years
(e85)	PS, MC	BOX	158	96.1% after 4 years
(e86)	PS, MC	STAR (318), Buechel-Pappas (92), AES (69), HINTEGRA (29), Mobility (23)	531	78% after 5 years, 62% after 10 years
(e87)	PS, MC	STAR (322), Mobility (132), AES (115), Buechel-Pappas (109), CCI (66), HINTEGRA (36)	780	81% after 5 years, 69% after 10 years
(e11)	RS, SC	STAR	123	86% (patients with preoperative deformity up to 10°), 75% (patients with preoperative deformity 10 to 30°) after 5 years
(e88)	PS, MC	Agility (117), STAR (45), Mobility (29), Ramses (11)	202	86% after 5 years
(e90)	RS, SC	Mayo	204	79% after 5 years, 65% after 10 years, 61% after 15 years
(e14)	RS, SC	Agility	132	86% after 9 years, 63% after 11 years
(e92)	PS, SC	STAR	100	85.7% (patients under 50) and 91.6% (patients over 50) after 5 years, 75% (patients under 50) and 80.6% (patients over 50) after 10 years
(e96)	PS, MC	BOX	189	97% after 4 years
(e16)	PS, SC	INBONE (211), STAR (122), Salto-Talaris (71)	404	90% and 97.6% after 3.2 years with and without arthrodesis of the hindfoot
(21)	RS, MC	Salto (91), HINTEGRA (39), AES (20), Coppelia (17), STAR (11), Ramses (4), Akile (1)	183	86% (88.4% high-volume centers; 84.9% low-volume centers) after 5 years
(e22)	PS, SC	Mobility	240	97.7% after 4 years
(e102)	RS, SC	Salto	401	86.6% (all patients), 85.1% (posttraumatic osteoarthritis), 95.6% (rheumatoid arthritis), 87.9% (patients under 55) after 5 years
(e25)	PS, MC	AES (298), STAR (217)	515	83% after 5 years
(e26)	RS, SC	Agility	306	80% (89% in patients over 54) after 5 years
(e108)	PS, SC	STAR	200	92.7% after 5 years
(e109)	RS, MC	Salto	109	97.5% after 2 years
(e111)	PS, SC	STAR	200	93.3% after 5 years, 80.3% after 10 years
(e112)	PS, SC	Buechel-Pappas (100), STAR (100)	200	79% (Buechel-Pappas) and 95% (STAR) after 6 years
(e30)	PS, SC	Mobility	100	97% after 3 years, 93.6% after 4 years

\*Clinical studies (total ankle replacement) with at least 100 patients were included.

AES: Ankle Evolutive System; BOX: Bologna-Oxford; MC: multicenter; PS: prospective; RS: retrospective; STAR: Scandinavian Total Ankle Replacement; TPR: Thomson, Prichard and Richard; SC: single-center

Trabecular Metal™) can be used to bridge the defect (37, 38, e113–e118). The alternative to converting to ankle arthrodesis is revision ankle arthroplasty (16, 17, e35, e119–e121). If possible, an implant type for which special revision components are available, e.g. a thicker metal plate for tibial components and larger weightbearing area and improved fixation for talar components, should be used. Revision surgery can be performed as one-stage or two-stage procedure. In the two-stage procedure, the goal of the first surgery is to address the bone defect. After bone integration of the autograft is achieved, the revision components can be implanted in a second surgery (eFigure 2).

### Conclusion

There is no gold standard treatment for advanced ankle osteoarthritis. Both, ankle arthrodesis and total ankle replacement are important treatment options in patients with end-stage ankle osteoarthritis. Attaining satisfactory intermediate-term and long-term postoperative outcomes in patients who have undergone total ankle replacement requires thorough preoperative examination and planning, taking careful account of all relative and absolute contraindications, with corresponding patient selection. If modern ankle implant designs are used, 10-year success rates of between 70 and 90% can be achieved.

### Conflict of interest statement

The authors declare that no conflict of interest exists.

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TABLE 3

Most common causes of failure of total hip, knee, and ankle replacement (given in % according to Sadoghi et al. [34])

Cause of failure	Total hip replacement	Total knee replacement	Total ankle replacement
Aseptic loosening	55.2	29.8	38
Luxation/instability	11.8	6.2	8.5
Septic loosening	7.5	14.8	9.8
Periprosthetic fracture	6	3	2
Pathological wear	4.2	8.2	8
Persistent pain	3.7	9.5	12
Implant failure	2.5	4.7	5.3
Technical error	3.8	4.6	4.6

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REVIEW ARTICLE

# Total Ankle Replacement

Indications, Implant Designs, and Results

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**eTABLE 1**

**Etiology of advanced ankle osteoarthritis, based on a selection of clinical and epidemiological studies\***

Study	Study type	Patients (ankles)	Etiology of ankle osteoarthritis, % (absolute values)		
			Primary	Secondary	Posttraumatic
(e6)	RS, SC, clinical	45 (51)	25.5% (13)	54.9% (28) – Rheumatism: 54.9% (28)	19.6% (10)
(6)	PS, SC, clinical	684 (722)	9.5% (69)	11.4% (82)	79.1% (571)
(e7)	PS, SC, clinical	47 (50)	6.0% (3)	8.0% (4) – Rheumatism: 4.0% (2) – Tuberculosis: 2.0% (1) – Hemophilia: 2.0% (1)	86.0% (43) – State following fracture: 50.0% (25) – State following ligament injuries: 36.0% (18)
(e8)	PS, SC, clinical	49 (50)	16.0% (8)	18.0% (9) – Rheumatism: 14.0% (7) – Necrosis of the talus: 4.0% (2)	66.0% (33)
(e9)	PS, MC register, clinical	245 (257)	20.6% (53)	55.3% (142)	24.1% (62) – State following fracture: 22.1% (57) – State following ligament injuries: 1.9% (5)
(e10)	PS, SC, clinical	80 (83)	33.7% (28)	25.3% (21) – Rheumatism: 25.3% (21)	41.0% (34)
(e11)	RS, SC, clinical	111 (123)	52.8% (65)	18.7% (23)	28.5% (35)
(e12)	RS, SC, clinical	61 (62)	19.4% (12)	4.8% (3) – Clubfoot: 3.2% (2) – Postpolio: 1.6% (1)	75.8% (47)
(e13)	RS, SC, clinical	45 (52)	50.0% (26)	26.9% (14)	23.1% (12)
(e14)	RS, SC, clinical	126 (132)	28.8% (38)	25.0 (33) – Rheumatism: 23.5% (31) – Postinfection: 1.5% (2)	46.2% (61)
(e15)	PS, SC, clinical	43 (50)	54.0% (27)	32.0% (16)	14.0% (7)
(e16)	PS, SC, clinical	396 (404)	16.6% (67)	13.6% (55)	69.8% (282)
(e17)	PS, SC, clinical	80 (84)	25.0% (21)	19.0% (16) – Rheumatism: 17.9% (15) – Hemochromatosis: 1.2% (1)	56.0% (47) – State following fracture: 39.3% (33) – State following ligament injuries: 16.7% (14)
(e18)	PS, SC, clinical	82 (82)	34.2% (28)	13.4% (11)	52.4% (43)
(e19)	RS, SC, clinical	95 (100)	26.0% (26)	29.0% (29) – Rheumatism: 26.0% (26) – Postinfection: 2.0% (2) – Psoriasis: 1.0% (1)	45.0% (45)
(e20)	PS, SC, clinical	229 (229)	13.8% (32)	4.0% (9)	82.2% (188)
(e21)	PS, SC, clinical	106 (106)	52.8% (56)	20.8% (22)	26.4% (28)
(e22)	PS, SC, clinical	233 (240)	30.8% (74)	17.9% (43) – Rheumatism: 15.0% (36) – Hemochromatosis: 2.9% (7)	51.3% (123)
(3)	PS, SC, epidemiological	639 (639)	7.2% (46)	23.2% (148) – Rheumatism: 11.9% (76) – Neuropathy: 4.9% (31) – Hemophilia: 1.9% (12) – Postinfection: 1.6% (10) – Gout: 0.8% (5)	69.6% (445) – State following fracture: 46.6% (298) – State following ligament injuries: 19.7% (126) – Traumatic osteochondral lesions: 3.3% (21)
(7)	PS, MC, clinical	593 (593)	26.5% (157)	15.3% (91)	58.2% (345)
(e23)	RS, SC, clinical	100 (100)	30.0% (30)	44.0% (44)	26.0% (26)
(e24)	RS, MC, clinical	478 (489)	15.3% (75)	25.8% (126)	58.9% (228)
(e25)	PS, MC register, clinical	515 (515)	19.2% (99)	59.2% (305)	21.6% (111)
(e26)	RS, SC, clinical	303 (306)	25.2% (77)	10.1% (31)	64.7% (198)
(e27)	RS, SC, clinical	103 (103)	71.8% (74)	18.5% (19)	9.7% (10)

Study	Study type	Patients (ankles)	Etiology of ankle osteoarthritis, % (absolute values)		
			Primary	Secondary	Posttraumatic
(e28)	PS, SC, clinical	65 (68)	13.2% (9)	16.2% (11) – Rheumatism: 7.4% (5) – Hemochromatosis: 4.4% (3) – Psoriasis: 1.5% (1) – Lupus erythematosus: 1.5% (1) – Sclerodermia: 1.5% (1)	70.6% (48) – State following fracture: 60.3% (41) – State following ligament injuries: 10.3% (7)
(e3)	RS, SC, epidemiological	390 (406)	8.9% (36)	12.8% (52) – Rheumatism: 5.4% (22) – Hemochromatosis: 2.7% (11) – Hemophilia: 1.5% (6) – Clubfoot: 1.0% (4) – Avascular necrosis of the talus: 0.7% (3) – Osteochondral lesions: 0.7% (3) – Postinfection: 0.7% (3)	78.3% (318) – State following fracture: 62.3% (253) – State following ligament injuries: 16.0% (65)
(e29)	PS, SC, clinical	66 (66)	0.0% (0)	10.6% (7) – Necrosis of the talus: 4.5% (3) – Hemophilia: 3.0% (2) – Rheumatism: 1.5% (1) – Hemochromatosis: 1.5% (1)	89.4% (59) – State following fracture: 72.7% (48) – State following ligament injuries: 16.7% (11)
(e4)	RS, SC, clinical	226 (233)	5.6% (13)	23.2% (54) – Flatfoot: 8.2% (19) – Rheumatism: 7.3% (17) – Clubfoot: 3.4% (8) – Osteochondral lesion: 1.7% (2) – Postinfection: 1.3% (1.3) – Charcot–Marie–Tooth disease: 0.9% (2) – Hemophilia: 0.4% (1)	71.2% (166) – State following fracture: 59.2% (138) – State following ligament injuries: 12.0% (28)
(e30)	PS, SC, clinical	96 (100)	64.0% (64)	27.0% (27) – Rheumatism: 27.0% (27)	9.0% (9) – State following fracture: 9.0% (9)
(e31)	RS, SC, clinical	90 (99)	40.4% (40)	12.1% (12)	47.5% (47)
<b>Total</b>		6218 (6389)	21.5% (1373)	22.6% (1441)	56.0% (3575)

\*Clinical (total ankle replacement) and epidemiological (etiology of ankle osteoarthritis) studies with at least 50 patients were included.  
MC: multicenter; PS: prospective; RS: retrospective; SC: single-center



**eTABLE 2**

**Classification of current ankle implant types**

	Surgical access	Inlay type	Replaced surfaces	Internal implant surfaces	Inlay materials	Sulcus type	Surface morphology
<b>AAA</b>	Anterior	Mobile	Superior	HA	UHMWPE	None	Trapezoidal
<b>AES</b>	Anterior	Mobile	Superior	HA	UHMWPE	Deep	Trapezoidal
<b>Agility</b>	Anterior	Fixed	Superior/medial/lateral	Titanium	UHMWPE	None	Trapezoidal
<b>BOX</b>	Anterior	Mobile	Superior	HA	UHMWPE	Normal	Ellipsoidal
<b>Buechel-Pappas</b>	Anterior	Mobile	Superior	Titanium	UHMWPE	Deep	Ellipsoidal
<b>ESKA</b>	Lateral/medial	Fixed	Superior	Titanium	UHMWPE	Normal	Ellipsoidal
<b>INBONE</b>	Anterior	Fixed	Superior	Titanium	UHMWPE	Normal	Spheroidal
<b>HINTEGRA</b>	Anterior	Mobile	Superior/medial	HA	UHMWPE	None	Conical
<b>Mobility</b>	Anterior	Mobile	Superior	Titanium	UHMWPE	Deep	Trapezoidal
<b>Ramses</b>	Anterior	Mobile	Superior	HA	UHMWPE	None	Ellipsoidal
<b>Salto</b>	Anterior	Mobile	Superior/medial	HA	UHMWPE	Normal	Conical
<b>STAR</b>	Anterior	Mobile	Superior	HA	UHMWPE	None	Cylindrical
<b>TNK</b>	Anterior	Fixed	Superior	HA	UHMWPE, ceramic	None	Cylindrical
<b>TM Total Ankle</b>	Lateral	Fixed	Superior	Porous metal	Highly cross-linked UHMWPE	None	Conical

AAA: Alpha Ankle Arthroplasty; AES: Ankle Evolutive System; BOX: Bologna–Oxford; HA: hydroxyapatite; STAR: Scandinavian Total Ankle Replacement; TM: Trabecular Metal; UHMWPE: ultra-high-molecular-weight polyethylene

**eTABLE 3**

**Additional procedures in patients with concomitant valgus or varus hindfoot deformity**

Patients with valgus hindfoot deformity	
Supramalleolar valgus deformity	Supramalleolar tibial osteotomy: – Medial closing tibial osteotomy (e43–e45)
Isolated valgus defective heel position	Corrective calcaneal osteotomy: – Medial sliding calcaneal osteotomy (e46)
Flexible pes planovalgus et abductus deformity	Corrective calcaneal osteotomy: – Lateral sliding calcaneal osteotomy (e47)  Tendon transfer: – Flexor digitorum longus to tibialis posterior (e48)
Rigid pes planovalgus et abductus deformity	Corrective arthrodesis of the hindfoot:* – Subtalar arthrodesis (e49) – Triple arthrodesis with or without calcaneocuboidal joint (e50)
Medial instability	Medial ligament stabilization: – Anatomical reconstruction with transosseous sutures (e51) – Reconstruction with tendon autograft (e52)
Patients with varus hindfoot deformity	
Supramalleolar varus deformity	Supramalleolar tibial osteotomy: – Medial opening tibial osteotomy (with supramalleolar deformity <10°) (e44, e45, e53) – Lateral closing tibial osteotomy (with supramalleolar deformity >10°) (e44, e45, e53)
Flexible varus abnormal heel position	Corrective calcaneal osteotomy: – Dwyer calcaneal osteotomy (e54) – Z-shaped calcaneal osteotomy (e55)
Rigid inframalleolar varus alignment	Corrective arthrodesis of the hindfoot: – Subtalar arthrodesis (e49)
Lateral instability	Lateral ligament stabilization: – Anatomical reconstruction with transosseous stitches (e56) – Reconstruction with tendon autograft (e57)

\*Depends on extent of deformity and degenerative changes

**eTABLE 4**

**Clinical outcomes following total ankle replacement: probability of survival of implant components and postoperative range of ankle motion\***

Study	Study type	Implant type	No. of implants	Probability of survival of implant components	Mean follow-up time	Postoperative range of motion
(e71)	RS, SC	Buechel–Pappas	35	97% after 5 years	5 years (3 to 150 months)	N/A
(e72)	RS, SC	Agility	207	76% after 9 years	N/A	N/A
(e73)	RS, SC	AES	93	90% after 5 years	42 months (13 to 73 months)	N/A
(e6)	RS, SC	STAR	51	70% after 5 years	52 months (36 to 97 months)	Total 28° (10 to 55°)
(22)	RS, SC	HINTEGRA	123	93% after 6 years	67.7 ± 27.0 months (29 to 126 months)	Total 35.3° ± 8.1°
(e34)	PS, SC	HINTEGRA	52	91% after 5 years, 78% after 8 years	5 years (2 to 10 years)	Total 38° ± 9°
(6)	PS, SC	HINTEGRA	722	94% after 5 years, 84% after 10 years	6.3 ± 2.9 years (2 to 12.2 years)	N/A
(e74)	RS, MC	Salto (388), AES (173), HINTEGRA (22), STAR (9)	592	88% after 71 months	Min. 1 year	N/A
(e75)	PS, SC	BOX	62	91.9% after 42.5 months	42.5 months (24 to 71 months)	DF 8.4° ± 4.8° (0 to 20°); PF 17.1° ± 8.3° (0 to 30°)
(e76)	RS, SC	Salto	98	98% after 5 years	35 months (24 to 68 months)	Total 28.3° ± 7°
(e77)	RS, SC	Salto	98	85% after 10 years	8.9 years (6.8 to 11 years)	DF 8.6° ± 5.3° (–5 to 20°); PF 18.1° ± 7.8° (5 to 40°)
(e78)	PS, SC	STAR	77	70.7 after 10 years, 45.6% after 14 years	12.4 years (10.8 to 14.9 years)	Total 22.8° ± 3.5°
(e8)	PS, SC	Buechel–Pappas	50	93.5% after 10 years	5 years (2 to 10 years)	Total 28° (12 to 46°)
(e79)	PS, SC	Buechel–Pappas (normal sulcus 40; deep sulcus 75)	115	74.2% (normal sulcus) after 20 years, 92% (deep Sulkus) after 12 years	12 years (2 to 10 years, normal sulcus), 5 years (2 to 12 years, deep sulcus)	Total 25° (10 to 47°, normal sulcus), total 29° (10 to 50°, deep sulcus)
(e80)	PS, SC	BOX	20	N/A	12 months (7 to 14 months)	Total 28.8° ± 11.3° (10 to 50°)
(e81)	RS, SC	Agility	42	62% after 9 years	8 years (0.5 to 11 years)	
(e82)	RS, SC	Buechel–Pappas	30	87.6% after 5 years	5.1 ± 4 years (1 to 13 years)	DF 5°; PF 30°
(e83)	PS, SC	LCS (19), Buechel–Pappas (74)	93	84% after 8 years	7.2 years (0.4 to 16.3 years)	DF 7.1° (5.8 to 8.4°); PF 24.8° (22.6 to 27.2°)
(e9)	PS, MC	STAR (216), TPR (32), HINTEGRA (6), AES (3)	257	89% after 5 years, 76% after 10 years	4 years (5 days to 12 years)	N/A
(e84)	PS, MC	BOX	51	97.2% after 3 years	30 months (24 to 48 months)	Total 27.4° (16 to 53°)
(e85)	PS, MC	BOX	158	96.1% after 4 years	17 months (6 to 48 months)	Total 26.5° (14 to 53°)
(e86)	PS, MC	STAR (318), Buechel–Pappas (92), AES (69), HINTEGRA (29), Mobility (23)	531	78% after 5 years, 62% after 10 years	1 to 11 years	N/A

Study	Study type	Implant type	No. of implants	Probability of survival of implant components	Mean follow-up time	Postoperative range of motion
(e87)	PS, MC	STAR (322), Mobility (132), AES (115), Buechel-Pappas (109), CCI (66), HINTEGRA (36)	780	81% after 5 years, 69% after 10 years	10 years	N/A
(e11)	RS, SC	STAR	123	86% (patients with preoperative deformity up to 10°) and 75% (patients with preoperative deformity 10 to 30°) after 5 years	4 years (2 to 8 years)	N/A
(e88)	PS, MC	Agility (117), STAR (45), Mobility (29), Ramses (11)	202	86% after 5 years	28 to 75 months	N/A
(e12)	RS, SC	Agility	65	91% after 1 year, 70% after 3 years, 67% after 5 years	3.3 years (2 to 5.9 years)	N/A
(e89)	RS, SC	TPR	33	85% after 10 years	10 to 23 years	N/A
(e13)	RS, SC	STAR	52	90% after 5 years, 84% after 8 years	80 months (60 to 110 months)	Total 23° ± 12° (0 to 55°)
(e90)	RS, SC	Mayo	204	79% after 5 years, 65% after 10 years, 61% after 15 years	9 years (2 to 17 years)	k. A.
(e14)	RS, SC	Agility	132	86% after 9 years, 63% after 11 years	9 years	DF 0° (-24 to 16°); PF 19° (-1 to 36°)
(e58)	PS, SC	STAR	28	70% after 12 years	1 to 12 years	N/A
(e91)	PS, SC	STAR	52	72.7% (primary osteoarthritis) and 75.5% (rheumatoid arthritis) after 14 years	9 years (6 to 14 years)	N/A
(e92)	PS, SC	STAR	100	85.7% (patients under 50) and 91.6% (patients over 50) after 5 years, 75% (patients under 50) and 80.6% (patients over 50) after 10 years	6.8 years (1 to 15 years)	N/A
(e93)	PS, SC	STAR (33 cemented, 25 uncemented)	58	70% (cemented) and 95.4% (uncemented) after 12 years	9.4 ± 2.7 years	N/A
(e94)	PS, SC	AES	38	79% after 2 years	28 months (2 to 70 months)	N/A
(e95)	PS, SC	LCS (19), Buechel-Pappas (74)	93	80% after 15 years	14.8 years (10.7 to 22.8 years)	N/A
(e96)	PS, MC	BOX	189	97% after 4 years	21 months	Total 14 to 53°
(e16)	PS, SC	INBONE (211), STAR (122), Salto-Talaris (71)	404	90% and 97.6% after 3.2 years with and without arthrodesis of the hindfoot	3.2 years (2 to 6 years)	N/A
(e17)	PS, SC	STAR	84	96% after 5 years, 90% after 10 years	9.1 years (2.6 to 11 years)	DF 4.5°; PF 34.7°
(e97)	RS, SC	AES	38	94.7% after 6 years	57.8 months (48 to 80 months)	N/A
(e98)	RS, SC	TNK	27	77% after 14.1 years	72 months (15 to 169 months)	DF 7.5° (0 to 20°); PF 8.5° (-10 to 20°)
(e99)	RS, SC	Salto	75	98% after 3.6 years	43 months (27 to 73 months)	DF 8.7° ± 5.6°; PF 29° ± 7°
(21)	RS, MC	Salto (91), HINTEGRA (39), AES (20), Coppelia (17), STAR (11), Ramses (4), Akile (1)	183	86% (high-volume sites 88.4%; low-volume sites 84.9%) after 5 years	39 ± 29 months (6 to 132 months)	N/A

Study	Study type	Implant type	No. of implants	Probability of survival of implant components	Mean follow-up time	Postoperative range of motion
(e100)	RS, MC	STAR	59	88% after 3 years	36 months (12 to 65 months)	DF 10.2° ± 6.3°; PF 11.3° ± 7.9°
(e22)	PS, SC	Mobility	240	97.7% after 4 years	32.8 ± 15.3 months (12 to 63 months)	DF 8.3° ± 5.3°; PF 13.6° ± 6.4°
(e101)	RS, SC	Buechel–Pappas	28	93% after 8.3 years	8.3 years (5 to 12.2 years)	Total 23° (8 to 40°)
(e102)	RS, SC	Salto	401	86.6% (all patients), 85.1% (posttraumatic osteoarthritis), 95.6% (rheumatoid arthritis), 87.9% (patients under 55) after 5 years	29 months (1 to 84 months)	Total 33.1° ± 13.6°
(e103)	PS, SC	TPR (20), STAR (19)	39	87% (TPR) after 12 years, 94.3% (STAR) after 6 years	8.6 years (TPR: 3 to 13 years), 3.1 years (STAR: 1 to 6 years)	Total (TPR) 37°. Total (STAR) 33.5°
(e104)	PS, SC	Salto Talaris	75	96% after 2.8 years	2.8 years (2 to 4.5 years)	N/A
(e25)	PS, MC	AES (298), STAR (217)	515	83% after 5 years	3.2 years (0.1 to 9.6 years)	N/A
(e26)	RS, SC	Agility	306	80% (89% in patients over 54) after 5 years	33 ± 18 months (4 to 75 months)	N/A
(e105)	PS, MC	Mobility	88	89.6% after 3 years, 88.4% after 4 years	40 months (30 to 60 months)	N/A
(e106)	RS, SC	Mobility	58	84% after 4 years	32 months (14 to 49 months)	N/A
(e107)	RS, SC	AES (16), Salto (4), New-Jersey (1)	21	91% after 3 years, 57% after 5 years	38 ± 26 months	N/A
(e108)	PS, SC	STAR	200	92.7% after 5 years	46 months (24 to 101 months)	N/A
(e109)	RS, MC	Salto	109	97.5% after 2 years	21.7 months (12 to 65 months)	Total 32°
(e110)	RS, SC	HINTEGRA	16	66.7% after 5 years	61.8 months (7 to 116 months)	Total 23.7° (12.0 to 47.5°)
(e111)	PS, SC	STAR	200	93.3% after 5 years, 80.3% after 10 years	88 months (60 to 156 months)	N/A
(e112)	PS, SC	Buechel–Pappas (100), STAR (100)	200	79% (Buechel–Pappas) and 95% (STAR) after 6 years	Min. 36 months	N/A
(e30)	PS, SC	Mobility	100	97% after 3 years, 93.6% after 4 years	43 months (4 to 63 months)	DF 7.5° (-5 to 22°), PF 14° (1 to 41°)

\*All available clinical studies (total ankle replacement) were included.

AES: Ankle Evolutive System; BOX: Bologna–Oxford; DF: dorsiflexion; N/A: information not available; LCS: low-contact stress; MC: multicenter; PF: plantar flexion; PS: prospective; RS: retrospective; STAR: Scandinavian Total Ankle Replacement; TPR: Thomson, Prichard and Richard; SC: single-center







